

NO 1169

Objective:

- Measure biomarkers of exposure in adult smokers switching from conventional cigarettes to ACCORD products for a period of 8 days,
- Determine if CO in exhalate and Carboxy-hemoglobin are at least 80% lower in smokers of ACCORD and OASIS compared to smokers of conventional cigarettes (3 – 6.9 MG TAR)

Deliverable(s):	Final report, publications and posters
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Reason for Doing this Work:	Development of standard approaches for new products development
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Program Area: ☐ Cancer ☐ CVD ☐ COPD ☐ Repro ☐ ETS ☐ Smoking Behavior ☒ Clinical Testing ☐ Communication

(Select One) ☐ Acceptability Assessment ☐ R.H. Evaluation ☐ R.H. Guidance ☐ Non-Clinical Testing ☐ Non-Clinical Research

Project Leader:	Hans J Roethig
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Tactics and Milestones:		Target Date:
Draft report		1/31/02
Results Ames testing		5/31/02
Final report		6/30/02
Publications:		
ETS		1/31/02
Cigarette burn time		5/31/02
Study design and smoking behavior		9/30/02
Exposure biomarkers		10/31/02
Potential harm biomarkers		10/31/02
Mutagenicity		12/31/02
Internal Resource Allocation (WSA, INBIFO) Franz Tewes 80 hrs, Robin Kinser 100 hrs, Bettie Nelson 150 hrs, Raymond Lau 35 hrs		External Resource Allocation (other PMUSA, external vendors)

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